WE CLAIM

A stent, comprising: **-1**.

a helical structure having a plurality of coils, said structure having a longitudinal axis and said coils having a pitch, said structure having an internal longitudinal passage wherein said structure is made from a filament having a crosssection and an outer surface, said filament comprising:

a soft flexible elongated member having an outer surface; and

a bioabsorbable or biodegradable polymeric outer coating on the outer surface of the member;

wherein, the polymeric coating has sufficient mechanical integrity to effectively maintain the flexible member in a helical configuration, until the coating has sufficiently been degraded or absorbed in vivo to effectively convert the helical structure back into a soft, elongated member.

- 2. The stent of claim 1, wherein the coating comprises a melt polymer.
- The stent of claim 1, wherein the coating comprises a solution polymer. **∕**3.
- The stent of claim 1 wherein the filament comprises a surgical suture. 4.

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- t 5. The stent of claim 4, wherein the suture comprises a monofilament.
 - 76. The stent of claim 4, wherein the suture comprises a multifilament.
 - 7. The stent of claim 4, wherein the suture comprises a non-absorbable suture.
 - 103 8. The stent of claim 4 wherein the suture comprises an absorbable suture.
 - The stent of claim 1, wherein the coating comprises a polymer made from monomers selected from the group consisting of lactide, glycolide, para-dioxanone, caprolactone, and trimethylene carbonate, blends thereof and copolymers thereof.
- 10. The stent of claim 1, wherein the polymer of the coating has a glass transition temperature above 55 C.
- 11. The stent of claim 1 wherein the polymer of the coating has a glass transition temperature above 120 C.
- The stent of claim 1, wherein the polymeric coating comprise a polymer selected from the group consisting of polyacrylamides, polyethylene glycols, polyethylene oxide, vinyl alcohols, and poly(N-vinyl pyrrolidone)s.
- # (3) 13. The stent of claim 1, wherein the polymeric coating additionally comprises polyamide.
- 14. A biodegradable filament, the filament comprising: ETH-1554

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an elongated, flexible member having a cross-section, and an outer surface; and,

a polymeric coating on said outer surface, said coating comprising a biodegradable or bioabsorbable polymer,

wherein, the polymeric coating has sufficient mechanical integrity to effectively maintain the flexible member in a substantially fixed configuration, until the coating has sufficiently been degraded or absorbed in vivo to effectively convert the structure back into a soft, elongated member.

- The filament of claim 14, wherein the coating comprises a melt polymer. ∠15.
- The filament of claim 14, wherein the coating comprises a solution *¥*6. polymer.
 - The filament of claim 14, wherein the filament comprises a surgical suture. *¥*7.
 - The filament of claim 17, wherein the suture comprises a monofilament. 63 18.
 - 19. The filament of claim 17, wherein the suture comprises a multifilament.
- *2*0. The filament of claim 17, wherein the suture comprises a non-absorbable suture.

The filament of claim 17 wherein the suture comprises an absorbable

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providing a stent, said stent comprising:

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suture.

- 72. The filament of claim 14, wherein the coating comprises a polymer made from monomers selected from the group consisting of lactide, glycolide, paradioxanone, caprolactone, and trimethylene carbonate, blends thereof and copolymers thereof.
- 23. The filament of claim 14, wherein the polymer of the coating has a glass transition temperature above 55 C.
- 24. The filament of claim 14 wherein the polymer of the coating has a glass transition temperature above 120 C.
- 25. The filament of claim 14, wherein the polymeric coating comprise a polymer selected from the group consisting of polyacrylamides, polyethylene glycols, polyethylene oxide, vinyl alochols, and poly(N-vinyl pyrrolidone)s.
- 26. The filament of claim 14, wherein the polymeric coating additionally comprises polyamide.
- √27. A method of maintaining a passageway of a body lumen substantially open, comprising the steps of:

a helical structure having a plurality of coils, said structure having a longitudinal axis and a longitudinal passage, and said coils having a pitch, wherein said structure is made from a fiber, said fiber having a cross-section and said filament comprising:

an elongated flexible, filament member, having an external surface and a crosssection; and,

a polymeric outer coating on the surface of the member, wherein, the polymeric coating has sufficient mechanical integrity to effectively maintain the flexible member in a helical configuration; and,

implanting said stent in a body lumen and maintaining the stent in the body lumen for a sufficient period of time to effectively maintain the passageway of the lumen substantially open for a desired period of time until the exterior coating softens, thereby converting the stent structure into a soft, flexible filamentary structure.